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                 chemical name field
NEWS
        OCT 06
                 Increase your retrieval consistency with new formats or
                 for Taiwanese application numbers in CA/CAplus.
NEWS
        OCT 21
                 CA/CAplus kind code changes for Chinese patents
                 increase consistency, save time
         OCT 22
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                 highlighting of terms when patent documents are
                 saved in .rtf format
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      6 OCT 28
                 INPADOCDB/INPAFAMDB: Enhancements to the US national
                 patent classification.
      7 NOV 03 New format for Korean patent application numbers in
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                 CA/CAplus increases consistency, saves time.
         NOV 04
                 Selected STN databases scheduled for removal on
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                 December 31, 2010
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                 Substance-Based Searching
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                 Search an additional 46,850 records with MEDLINE
                 backfile extension to 1946
         DEC 14 New PNK Field Allows More Precise Crossover among STN
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                 Patent Databases
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         DEC 18 ReaxysFile available on STN
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         DEC 21 CAS Learning Solutions -- a new online training experience
NEWS 15 DEC 22 Value-Added Indexing Improves Access to World Traditional
                 Medicine Patents in CAplus
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         JAN 24
                 The new and enhanced DPCI file on STN has been released
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         JAN 26 Improved Timeliness of CAS Indexing Adds Value to
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                 Updated MeSH vocabulary, new structured abstracts, and
NEWS 18
         JAN 26
                 other enhancements improve searching in STN reload of
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10593748.trn 02/22/2011 Page 1

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FILE 'HOME' ENTERED AT 09:46:19 ON 22 FEB 2011

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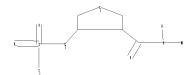
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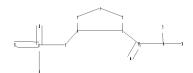
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```
chain nodes :
7  8  11  12  13  14  15  16  17  18
ring nodes :
1  2  3  4  5
chain bonds :
3-14  4-7  7-8  8-11  8-12  8-13  14-15  14-16  15-17  15-18
ring bonds :
1-2  1-5  2-3  3-4  4-5
exact/norm bonds :
1-2  1-5  2-3  3-4  3-14  4-5  4-7  7-8  8-11  8-12  8-13  14-15  14-16  15-17
15-18
isolated ring systems :
containing 1 :
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G1:0,S,N,CH2

G2:CH2,NH

G3:Cb, Cy, Hy, Ak, Ph

Match level :

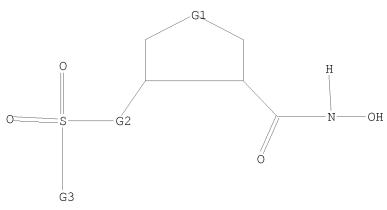
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 7:CLASS 8:CLASS 11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:CLASS 18:CLASS

# L1 STRUCTURE UPLOADED

=> d 11

L1 HAS NO ANSWERS

L1 STR



G1 O, S, N, CH2

G2 CH2, NH

G3 Cb, Cy, Hy, Ak, Ph

Structure attributes must be viewed using STN Express query preparation.

=> s 11

SAMPLE SEARCH INITIATED 09:47:18 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 29 TO ITERATE

100.0% PROCESSED 29 ITERATIONS 12 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*
BATCH \*\*COMPLETE\*\*

PROJECTED ITERATIONS: 257 TO 903 PROJECTED ANSWERS: 33 TO 447

L2 12 SEA SSS SAM L1

=> FIL HCAPLUS

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FILE COVERS 1907 - 22 Feb 2011 VOL 154 ISS 9
FILE LAST UPDATED: 21 Feb 2011 (20110221/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Oct 2010
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Oct 2010

HCAplus now includes complete International Patent Classification (IPC) reclassification data for the fourth quarter of 2010.

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> FIL REGISTRY
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SINCE FILE TOTAL
ENTRY SESSION
2.99 3.73

FULL ESTIMATED COST

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=> s l1 sss full FULL SEARCH INITIATED 09:47:40 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 589 TO ITERATE

100.0% PROCESSED 589 ITERATIONS SEARCH TIME: 00.00.01

229 ANSWERS

T.3 229 SEA SSS FUL L1

=> FIL HCAPLUS COST IN U.S. DOLLARS

ENTRY SESSION 200.59 FULL ESTIMATED COST 196.86

SINCE FILE

TOTAL

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FILE COVERS 1907 - 22 Feb 2011 VOL 154 ISS 9 FILE LAST UPDATED: 21 Feb 2011 (20110221/ED) REVISED CLASS FIELDS (/NCL) LAST RELOADED: Oct 2010 USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Oct 2010

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 13

L45 L3

=> s 14 and py<=2004 25160617 PY<=2004

 $L_5$ 3 L4 AND PY<=2004

=> d 15 ibib abs hitstr tot

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN ACCESSION NUMBER: 2002:539654 HCAPLUS DOCUMENT NUMBER: TITLE:

137:93692
Preparation of
(quinolinylmethoxyphenylsulfonylmethyl)-substituted
pyrrolidinecarboxamides and piperidinecarboxamides as
MMP, TNP, and/or aggreenase inhibitors
Xue, Chu-Biao; Decicco, Carl P.; Re, Xiaohua
Bristol-Myers Squibbl Company Patent Department, USA
PCT Int. Appl., 133 pp.
CODEN: PIXXD2
Patent

INVENTOR (S): PATENT ASSIGNEE(S):

DOCUMENT TYPE: LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION: English

PATENT NO. KIND DATE APPLICATION NO. DATE WO 2002055491 A2 20020718 WO 2002-US760 20020109 WO 2002055491 A3 20030123
W1 A2, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, BC, EE, ES, FI, GB, GD, GE, GB, GB, HR, HU, ID, IL, IN, IS, JP, KE, KG, RP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MM, MX, MZ, NO, NZ, CM, PH, FL, FT, RO, RU, SD, SE, SG, SI, SK, SI, TJ, TM, TH, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZM, ZW
RNI: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, LE, IT, LU, MC, NIL, FT, SE, TR, FF, FJ, CF, CG, CI, CM, GA, GN, GQ, GN, ML, MR, NE, SN, TD, TG
CA 2434044 A1 20020718 CA 2002-2434044 20020109

AU 2002246983 20020724 A1 AU 2002-246983 20020109 US 2002-43541 US 20030087890 A1 20030508 20020109

US 6642255 EP 1355648 20031104 20031029 B2 A2 EP 2002-714733 20020109

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR PRIORITY APPLN. INFO: US 2001-260957P P 20010111

WO 2002-US760 W 20020109

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT OTHER SOURCE(S): MARPAT 137:93692

OTHER SOURCE(S):

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) exhibited Ki values of ≤ 10 μM against MMP-1, 2, 3, 9, and 13. Thus, I are useful for the treatment of inflammatory disorders and thromboembolic disorder (no data).

IT 441297-34-3P
RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Usea) (MMP, TNF, and/or aggrecanase inhibitor; preparation of (quinoliny)methoxyphenyl.sulfonylmethyl)-substituted pyrrolidinecarboxamides and piperidinecarboxamides as MMP, TNF, and/or aggrecanase inhibitors;

RN 441297-34-3 HCAPLUS
CN 1-Pyrrolidinecarboxylic acid, 3-(hydroxyamino)carboxyl]-4-[[[4-[(2-methyl-4-quinolinyl)methoxyl]phenyl]sulfonyl]methyl]-, (3R, 48)-, 1,-dimethylethyl
ester, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CRN 441297-33-2 CMF C28 H33 N3 O7 S

Absolute stereochemistry.

PAGE 1-A

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

Title compds. I [wherein A = COR5, CO2H, CH2CO2H, CO2R6, CONHOH, CONHOR5, CONHOR6, N(OH)CHO, N(OH)COR5, SH, CH2SH, SONHRA, SN2H2Ra, PO3H2, or PO(OH)NHRA; ring B = 3-10 membered (hetero)cycle; Z = absent or (un)substituted (hetero)cycly; U = absent or O, NH, N(alkyl), CO, CO2, CCO, CONH, NHCO, CCO2, etc. X = absent or alkylene, alkenylene, or alkynylene; Y = absent or O, NH, N(alkyl), SOO-2, or CO; Za = (un)substituted (hetero)cyclyl; Rla and Rlb = independently H, alkyl, Ph, PhCH2, CH2OR3, or (un)substituted CH2NH2; or CR1aRlb = (hetero)cyclyl; R2 = Q or (un)substituted alkylene-Q, alkenylene-Q, or alkynylene-Q, Q-substituted alkoxy(alkyl), carbamoyl(alkyl), sulfamoyl(alkyl), etc.;

= H, alkyl, ORa, (un) substituted CH2NH2, or SOO-2Ra; R2b = H or alkyl; Q

H or (un)substituted (hetero)cycly1; R3 = Q1 or (un)substituted alkylene-Q1, alkenylene-Q1, or alkynylene-Q1, Q1-substituted alkoxy(alky1), carbamoy(alky1), sulfamoy(alky1), etc.; or C(R3)2 = (un)substituted (hetero)cycly1; Q1 = H or (un)substituted Ph, naphthy1,

or

heteroaryl; Ra = H, alkyl, Ph, or PhCH2; p = 0-2; R5 = (un)substituted
alkyl; R6 = phenyl(alkyl), naphthyl, cycloalkyl, alkylcarbonyloxy, etc.;
or pharmaceutically acceptable salt thereof] were prepared as matrix
metalloprotease (MMP), tumor necrosis factor (TNF), and aggrecanase
inhibitors. For example, the
3-(quinolinylmethoxyphenylsulfonylmethyl)-4piperidinecarboxamide (3R, 48)-II-2CFSCO2H was prepared in seventeen
steps starting from the reaction of N-benzyloxycarbonyl-β-alanine and
benzylbromide. Key steps include the cyclication of the 5-aminopentanal
intermediate and the addition of 4-mercaptophenol and
4-chloromethyl-2-methylquinoline+BCl. A number of invention compds.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 2-A

441297-33-2P 441297-37-6P 441297-40-1P 441297-35-4P 441297-38-7P IT 441297-36-5P 441297-39-8P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES

(MMP, TNF, and/or aggrecanase inhibitor; preparation of (MMP, TNF, and/or aggrecanase inhibitor; preparation of (quinoliny]methoxyphenylsulfonylmethyl)-substituted pyrrolidinecarboxamides and piperidinecarboxamides as MMF, TNF, and/or aggrecanase inhibitors)

RN 441297-33-2 BCAPLUS

RN 1-Pyrrolidinecarboxylic acid,
3-[(hydroxyamino)carbonyl]-4-[[[4-[(2-methyl4-quinolinyl)]methoxy]phenyl]sulfonyl]methyl]-, 1,1-dimethylethyl ester,

(3R,4S)- (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

441297-35-4 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R, 4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

441297-36-5 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R,48)-, 2,2-trifluoroacetate (1:2) (CA INDEX NAME)

CRN 441297-35-4 CMF C23 H25 N3 O5 S

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

CM 2

CRN 76-05-1 CMF C2 H F3 O2

RN 441297-37-6 BCAPLUS
CN 3-Pytrolidinecarboxamide,
N-hydroxy-1-(1-methylethyl)-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A

RN 441297-38-7 HCAPLUS
CN 3-Pyrrolidinecarboxamide,
N-hydroxy-1-(1-methylethyl)-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-,
2,2,2-trifluoroacetate (1:2) (CA INDEX NAME)

CM 1

CRN 441297-37-6 CMF C26 H31 N3 O5 S

PAGE 1-A

PAGE 2-A

CM 2 CRN 76-05-1 C2 H F3 O2

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN CRN 441297-39-8 CMF C26 H27 N3 O5 S (Continued)

Absolute stereochemistry.

PAGE 2-A

CM 2 CRN 76-05-1 CMF C2 H F3 O2

10593748.trn 02/22/2011

- ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) 441297-39-8 HCAPLUS 3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-1-(2-propyn-1-yl)-, (3R,4S)-(CA INDEX NAME)

Absolute stereochemistry.

PAGE 2-A

441297-40-1 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinoliny])methoxy]phenyl]sulfonyl]methyl]-1-(2-propyn-1-yl)-, (3R,4S)-, 2,2,2-trifluoroacetate (1:2) (CA INDEX NAME)

CM 1

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

101037-97-1 101038-00-9 101038-05-2 1101038-05-3 1101038-15-6 1101038-15-6 1101038-15-6 1101038-15-1 1101038-21-4 1101038-21-4 1101038-27-0 1101038-30-1 1101037-98-2 1101038-01-0 1101038-04-3 1101038-07-3 1101038-10-1 1101038-10-1 1101038-15-1 1101038-15-1 1101038-25-1 1101038-25-1 1101038-25-1 1101038-31-9 1101038-31-9 1101038-40-7 1101038-40-7 1101038-40-7 1101038-40-7 1101038-98-5 1101038-98-1 1101038-98-1 1101038-98-1 1101038-98-1 1101038-98-1 1101038-98-1 1101038-98-1 1101038-98-1 1101038-98-1 1101038-98-1 1101037-99-3 1101038-02-1 1101038-05-4 1101038-08-7 1101038-14-2 1101038-14-2 1101038-17-8 1101038-29-2 1101038-29-2 1101038-29-2 1101038-32-7 1101038-38-3 1101038-47-4 1101038-47-4 1101038-99-7 1101038-99-6 1101038-99-6 1101038-99-6 1101038-99-6 1101038-99-6 1101038-99-6 1101038-99-6 1101038-99-6 1101038-99-6 1101038-99-6 1101039-10-4 1101039-13-7 1101039-16-0 1101039-19-3 1101039-11-5 1101039-14-8 1101039-17-1 1101039-20-6 1101039-21-7 1101041-15-9 1101041-18-2 1101041-21-7 1101039-22-8 1101041-14-8 1101041-17-1 1101041-16-0 1101041-20-6 1101041-19-3 1101041-22-8 1101041-23-9 1101041-21-7 1101041-24-0 1101041-27-3 1101041-30-8 1101041-33-1 1101041-22-0 1101041-25-1 1101041-28-4 1101041-31-9 1101041-34-2 1101041-23-9 1101041-26-2 1101041-29-5 1101041-32-0 1101041-35-3 1101041-36-4 1101041-39-7 1101041-84-2 1101041-87-5 1101041-37-5 1101041-40-0 1101041-38-6 1101041-41-1 1101041-86-4 1101041-89-7 1101041-85-3 1101041-88-6 101041-90-0 101041-90-0 101041-98-3 101041-98-3 101041-98-3 101042-08-0 101042-08-0 101042-11-8 101042-11-8 101042-21-2 101042-23-2 101042-23-2 101042-23-2 101042-35-3 101042-35-6 101042-38-9 1101041-83-6 1101041-88-6 1101041-91-4 1101041-94-4 1101042-00-5 1101042-03-8 1101042-03-8 1101042-09-4 1101042-15-2 1101042-15-2 1101042-21-0 1101042-21-0 1101042-21-0 1101042-21-1 1101042-31-4 1101042-33-4 1101042-33-4 101041-80-7 101041-89-7 101041-95-5 101041-95-5 101042-01-6 101042-01-6 101042-01-7 101042-10-7 101042-13-7 101042-13-7 101042-13-7 101042-25-1 101042-25-1 101042-31-5 101042-31-5 101042-31-5 101042-31-5 101042-31-5 101042-31-5

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN 1101043-60-0 1101043-61-1 1101043-62-2 1101043-63-3 1101043-64-4 1101043-65-5 1101043-66-6 1101043-67-7 1101043-68-8 1101043-72-4 1101043-73-5 1101043-71-3 1101043-72-4 1101043-73-5 1101043-71-6 1101043-73-7 1101043-73-6 1101043-73-6 1101043-73-6 1101043-73-6 1101043-73-7 1101043-81-5 1101043-81-5 1101043-81-5 1101043-81-5 1101043-81-5 1101043-81-5 1101043-81-5 1101043-81-5 1101043-81-5 1101043-81-5 1101043-81-5 1101043-81-5 1101043-63-6 1101043-63-7 1101044-63-6 1101043-63-7 1101044-63-6 1101043-63-7 1101044-63-6 1101043-63-7 1101043-63-8 1101043-63-7 1101043-63-8 1101043-63-7 1101043-63-8 1101043-63-7 1101043-63-8 1101043-63-7 1101043-63-8 1101043-63-7 1101043-63-8 1101043-6 (Continued) 1101043-66-6 1101043-67-7 1101043-68-8 1101043-68-8 1101043-69-9 1101043-70-2 1101043-71-3 1101043-72-4 1101043-76-8 1101043-74-6 1101043-78-0 1101043-76-8 1101043-79-1 1101043-78-0 1101043-78-0 1101043-80-4 1101043-81-5 1101043-82-6 1101043-80-4 1101043-81-5 1101043-82-6 1101044-59-0 1101044-60-3 1101044-61-4 1101044-65-8 1101044-60-3 1101044-61-4 1101044-65-8 1101044-61-6 1101044-61-6 1101044-65-8 1101044-61-6 1101044-61-6 1101044-65-8 1101044-61-6 1101044-61-6 1101044-65-8 1101043-81-7 1101043-81 1101043-81-7 1101043-81-7 1101043-81-7 1101043-81-7 1101043-81-7 1101043-81-7 1101043-81-7 1101043-81-7 1101043-81-7 1101043-81 1101043-81 1101043-81 1101043-81 1101043-81 1101043-81 1101045-81 1101043-81 1101043-81 1101043-81 1101043-81 1101043-81 11010

1101037-98-2 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-(4-pyridinylmethoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101038-01-0 HCAPLUS
3-Pyrrolidinecarboxamide, 4-[[[4-[(2-chloro-6-methoxy-4-pyridiny])methoxy]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R, 4S)-(CA INDEX NAME)

Absolute stereochemistry.

 $\label{local-problem} $$101038-02-1$ HCAPLUS $$3-Pyrrolidinecarboxamide, $N-hydroxy-1-methyl-4-[[[4-(4-quinolinylmethoxy)phenyl]sulfonyl]methyl]-, $(3R,4S)-$ (CA INDEX NAME) $$$$ 

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101037-99-3 HCAPLUS
3-Fytrolidinecarboxamide, 4-[[[4-[(2,6-dimethyl-4-pyridinyl)methoxy]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)-(CA INDEX NAME)

1101038-00-9 HCAPLUS
3-Fyrrolidinecarboxamide, 4-[[4-[(2-chloro-6-methyl-4-pyridinyl)methoxy]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R, 4S)-(CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A

1101038-03-2 HCAPLUS

3-Pyrrolidinecarboxamide, N-hydroxy-1-methy1-4-[[[4-(5-quinolinylmethoxy)phenyl]sulfonyl]methy1]-, (3R,4S)- (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

 $\label{local-problem} \begin{array}{lll} 1101038-04-3 & \text{HCAPLUS} \\ 3-\text{Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-(6-quinoliny]methoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME) \\ \end{array}$ 

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 2-A

1101038-06-5 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

PAGE 2-A

1101038-07-6 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methoxy-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-1-methyl-, (3R,4S)- (CA INDEX

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101038-05-4 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-(5-isoquinolinylmethoxy)phenyl]sulfonyl]methyl]-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN NAME) (Continued)

Absolute stereochemistry.

PAGE 1-A

PAGE 2-A

1101038-08-7 HCAPLUS
3-Fyrrolidinecarboxamide, 4-[[[4-[(2-amino-4-quinoliny])methoxy]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)-(CA INDEX NAME)

PAGE 1-A

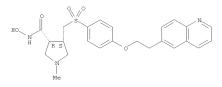
PAGE 2-A



1101038-09-8 BCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-[2-(4-quinolinyl)tethoxy]phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)



1101038-12-3 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[2-(5-isoquinolinyl)ethoxy]phenyl]sulfonyl]methyl]-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-13-4 HCAPLUS 3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-{[[4-[2-(2-methyl-4-quinolinyl)ethoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101038-10-1 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[4-[2-(5-quinolinyl)ethoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101038-14-5 HCAPLUS 3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[2-(2-methoxy-4-quinolinyl)ethoxy]phenyl]sulfonyl]methyl]-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-15-6 HCAPLUS
3-Pyrrolidinecarboxamide, 4-[[[4-[2-(2-amino-4-quinoliny])ethoxy]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R, 4S)-(CA INDEX NAME)

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101038-16-7 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4(phenoxymethyl)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

 $\label{local-problem} $$101038-17-8$$ BCAPLUS $$3-Pytrolidinecarboxamide, $4-[[4-[(3,5-dinethyl]phenoxy)methyl]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, $(3R,4S)-$$ (CA INDEX NAME) $$$ 

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

RN 1101038-20-3 HCAPLUS
CN 3-Pyrrolidinecarboxamide, 4-[[[4-[[(2,6-dimethyl-4-pyridinyl)oxy]methyl]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-,
(3R,4S)(CA INDEX NAME)

Absolute stereochemistry.

1101038-21-4 HCAPLUS 3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-[(4-pyridinyloxy)methyl]phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101038-18-9 HCAPLUS 
3-Pytrolidinecarboxamide, 4-[[[4-[(3,5-dichlorophenoxy)methyl]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, <math>(3R,4S)- (CA INDEX NAME)

1101038-19-0 HCAPLUS
3-Pyrrolidinecarboxamide, 4-[[4-[[3,5-bis(trifluoromethyl)phenoxy]methyl]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

RN 1101038-22-5 HCAPLUS
CN 3-Pyrrolidinecarboxamide, 4-[[[4-[[(2-chloro-6-methyl-4-pyridinyl)oxy]methyl]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-,
(3R,4S)(CA INDEX NAME) RN CN

Absolute stereochemistry.

1101038-23-6 HCAPLUS INDEX NAME NOT YET ASSIGNED

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101038-24-7 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[4-[(4-quinolinyloxy)methyl]phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

PAGE 1-A

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN

(Continued) PAGE 2-A

1101038-25-8 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-[(5-quinolinyloxy)methyl]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

PAGE 1-A

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 2-A

1101038-26-9 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[4-[(6-quinolinyloxy)methyl]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

1101038-27-0 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(5-isoquinolinyloxy)methyl]phenyl]sulfonyl]methyl]-1-methyl-, (3R, 4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A

1101038-28-1 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-[[(2-methyl-4-quinolinyl)oxy]methyl]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

1101038-29-2 HCAPLUS
3-Fyrrolidinecarboxamide, N-hydroxy-4-[[4-(1H-indol-5-yloxy)phenyl]sulfonyl]methyl]-1-methyl-, (3R,48)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

Absolute stereochemistry.

RN 1101038-33-8 HCAPLUS
CN 3-Pyrrolidinecarboxamide,
N-hydroxy-1-methyl-4-[[4-[[(1-methyl-1H-indol-5yl)oxy|methyl]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-34-9 HCAPLUS
3-Pyrrolidinecarboxamide, 4-[[[4-[[(1-ethyl-1H-indol-5-yl)oxy]methyl]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-35-0 HCAPLUS 3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-(1H-indol-5-

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

RN 1101038-30-5 HCAPLUS
CN 3-Pyrrolidinecarboxamide,
N-hydroxy-1-methyl-4-[[4-[(1-methyl-1H-indol-5yl)oxy]phenyl]sulfonyl]methyl]-, (3R, 4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-31-6 HCAPLUS
3-Pyrrolidinecarboxamide, 4-[[[4-[(1-ethyl-1H-indol-5-yl)oxy]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX

Absolute stereochemistry.

1101038-32-7 HCAPLUS

3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(1H-indol-5-yloxy)methyl]phenyl]sulfonyl]methyl]-1-methyl-, (3R,4S)- (CA INDEX NAME)

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) ylmethoxy)phenyl]sulfonyl]methyl]-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

RN 1101038-36-1 HCAPLUS
CN 3-Pyrrolidinecarboxamide,
N-hydroxy-1-methyl-4-[[4-([1-methyl-1H-indol-5yl)methoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-37-2 HCAPLUS
3-Pyrrolidinecarboxamide, 4-[[[4-(1H-benzimidazol-1-ylmethyl]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-38-3 HCAPLUS 3-Pyrrolidinecarboxamide, 4-[[[4-[(1-ethyl-1H-indol-5-

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) y1)methoxy]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

 $\label{local-equation} $$101038-39-4$ $$ HCAPLUS$ $$3-Pyrrolidinecarboxamide, $4-[[[4-[2-(1H-benzimidazol-1-yl)ethyl]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, $$(3R,4S)-$$(CA INDEX NAME)$ $$$ 

Absolute stereochemistry.

1101038-40-7 BCAPLUS
3-Pyrrolidinecarboxamide, 4-[[[4-(1H-benzotriazol-1-ylmethyl)phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) 1101038-43-0 HCAPLUS 3-Pyrrolidinecarboxamide, 4-[[[4-[(2,5-dimethyl-4-thiazolyl)methoxy]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)-(CA INDEX NAME)

Absolute stereochemistry

1101038-44-1 HCAPLUS
3-Pyrrolidinecarboxamide, 4-[[[4-[(4,5-dimethyl-2-thiazolyl)methoxy]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)-(CA INDEX NAME)

Absolute stereochemistry.

RN 1101038-45-2 HCAPLUS
CN 3-Pyrrolidinecarboxamide,
N-hydroxy-1-methyl-4-[[4-[[2-(1-methylethyl)-4thiazolyl]methoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

RN 1101038-46-3 HCAPLUS 10593748.trn

02/22/2011

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101038-41-8 HCAPLUS
3-Pyrrolidinecarboxamide, 4-[[[4-[2-(1H-benzotriazol-1-y1)ethyl]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-42-9 HCAPLUS
3-Pytrolidinecarboxamide, 4-[[4-[(2,4-dimethyl-5-thiazolyl)methoxy]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)-(CA INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) 3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-[(5-methyl-4-isoxazoly1)methoxy]phenyl]sulfonyl]methyl]-, (3R, 4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-47-4 HCAPLUS

N. 3-Pyrolidinecarboxamide,
N-hydroxy-1-methyl-4-[[(4-[[2-(1-methylethyl)-5-thiazolyl]methoxy]]henbyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

RN 1101038-48-5 HCAPLUS
CN 3-Pytrolidinecarboxamide,
4-[[[4-(2-butyn1-y)-loxy]phenyl]sulfonyl]methyl]N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-49-6 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-(2-pentyn-1-yloxy)phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-90-7 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[(phenylsulfonyl)methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-91-8 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[(4-methylphenyl)sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

RN 1101038-92-9 HCAPLUS CN 3-Pyrrolidinecarboxamide, N-hydroxy-4-[[(4-methoxyphenyl)sulfonyl]methyl]-1-methyl-, (3R,4S)- (CA INDEX NAME)

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued methylethoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-96-3 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-(2-methylpropoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-97-4 HCAPLUS
3-Pyrrolidinecarboxamide, 4-[[[4-(1,1-dimethylethoxy)phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

RN 1101038-98-5 HCAPLUS
CN 3-Pyrrolidinecarboxamide,
4-[[[4-(cyclopropyloxy)phenyl]sulfonyl]methyl]-Nhydroxy-1-methyl-, (3R,48)- (CA INDEX NAME)

10593748.trn 02/22/2011 Page 17 L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) Absolute stereochemistry.

RN 1101038-93-0 HCAPLUS
CN 3-Pytrolidinecarboxamide,
4-[[(4-ethoxyphenyl)sulfonyl]methyl]-N-hydroxy-1methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101038-94-1 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[(4-propoxyphenyl)sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME) CN

Absolute stereochemistry.

1101038-95-2 HCAPLUS 3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-(1-

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

RN 1101038-99-6 HCAPLUS
CN 3-Pyrrolidinecarboxamide,
4-[[[4-(cyclobutyloxy)phenyl]sulfonyl]methyl]-Nhydroxy-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

RN 1101039-00-2 HCAPLUS
CN 3-Pyrrolidinecarboxamide,
4-[[[4-(cyclopentyloxy)phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R, 4S)- (CA INDEX NAME)

Absolute stereochemistry.

RN 1101039-01-3 HCAPLUS CN 3-Pyrrolidinecarboxamide, 4-[[[4-(cyclohexyloxy)phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX NAME)

1101039-02-4 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[(4-phenoxyphenyl)sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

RN CN

 $\label{eq:continuous} \begin{tabular}{ll} $101039-03-5 & $HCAPLUS$ \\ $3-$Pyrrolidinecarboxamide, $4-[[[4-(3,5-dimethylphenoxy)phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, $(3R,4S)-dimethylphenoxy)phenyl]sulfonyllmethyl]-N-hydroxy-1-methyl-, $(3R,4S)-dimethylphenoxy)phenyllsulfonyllmethyllmethyllmethyllmethyllmethyllmethyllmethyllsulfonyllmethyll$ (CA

INDEX NAME)

Absolute stereochemistry.

 $\label{eq:continuous} \begin{tabular}{ll} $101039-04-6 & $HCAPLUS$ \\ $3-$Pyrrolidinecarboxamide, $4-[[[4-(3,5-dichlorophenoxy)phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, $(3R,4S)-dichlorophenoxy)phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, $(3R,4S)-dichlorophenoxy)phenyl]sulfonyl]methyl-, $(3R,4S)-dichlorophenoxy)phenyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]sulfonyl]$ 

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) 3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-(4-methylphenoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

RN 1101039-08-0 HCAPLUS
CN 3-Pytrolidinecarboxamide,
4-[[[4-(4-chlorophenox)]phenyl]sulfonyl]methyl]N-hydroxy-1-methyl-, (3R, 4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101039-09-1 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-(4-pyridinyloxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101039-10-4 HCAPLUS
3-Pyrrolidinecarboxamide, 4-[[4-[(2,6-dimethyl-4-pyridinyl)av]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX NAME)

10593748.trn 02/22/2011 Page 18

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN INDEX NAME) L5 (Continued)

Absolute stereochemistry.

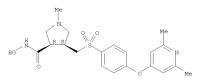
1101039-05-7 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-(3-methylphenoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

RN 1101039-06-8 HCAPLUS
CN 3-Pytrolidinecarboxamide,
4-[[[4-(3-chlorophenoxy)phenyl]sulfonyl]methyl]N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

RN 1101039-07-9 HCAPLUS

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) Absolute stereochemistry.



1101039-11-5 HCAPLUS INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

1101039-12-6 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-(5-quinolinylyphenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN L5 (Continued)

1101039-13-7 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-(6-quinolinyloxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101039-14-8 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-(5-isoquinolinyloxy)phenyl]sulfonyl]methyl]-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101039-15-9 HCAPLUS

3-Pyrrolidinecarboxamide, N-hydroxy-1-methy1-4-[[[4-[(2-methy1-4-quinoliny1)oxy]pheny1]sulfony1]methy1]-, (3R,4S)- (CA INDEX NAME (CA INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

Absolute stereochemistry.

1101039-19-3 HCAPLUS 
3-Pytrolidinecarboxamide, 4-[[4-[(3,5-dinethoxyphenyl)methoxy]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101039-16-0 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-(phenylmethoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

 $\label{local-problem} $$101039-17-1$$ BCAPLUS $$3-Fyrrolidine carboxamide, $4-[[[4-[(3,5-dimethylphenyl)methyl]+N-hydroxy-1-methyl-, (3R,4S)-$$ (CA INDEX NAME) $$$ 

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101039-20-6 HCAPLUS
3-Pyrrolidinecarboxamide, 4-[[4-[(3,5-dibromophenyl)methoxy]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S) (CA INDEX NAME)

Absolute stereochemistry.

1101039-21-7 HCAPLUS
3-Pyrrolidinecarboxamide, 4-[[4-[[3,5-bis(trifluoromethyl)phenyl]methoxy]phenyl]sulfonyl]methyl]-N-hydroxy-1-methyl-, (3R,4S)- (CA INDEX NAME)

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101039-22-8 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-methyl-4-[[[4-(2-pyridinylmethoxy)phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

1101041-14-8 HCAPLUS INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101041-15-9 HCAPLUS INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

1101041-16-0 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[(4-quinolinyloxy)methyl]phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101041-17-1 BCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[(5-quinolinyl)axy)methyl]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101041-18-2 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[(6-quinolinyloxy)methyl]phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) Absolute stereochemistry.

1101041-19-3 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[(5-isoquinolinyloxy)methyl]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101041-20-6 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[[(2-methyl-4-quinolinyl)oxy]methyl]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

10593748.trn

02/22/2011

Page 20

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101041-21-7 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(1H-indol-5-yloxy)phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

Absolute stereochemistry.

1101041-22-8 HCAPLUS

3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[(4-[(1-methyl-1H-indol-5-yl)oxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101041-26-2 HCAPLUS INDEX NAME NOT YET ASSIGNED

1101041-27-3 HCAPLUS

3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(1H-indol-5-ylmethoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101041-28-4 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[(1-methyl-1H-indol-5-yl)methoxy]phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101041-23-9 HCAPLUS
3-Furancarboxamide, 4-[[[4-[(1-ethyl-1H-indol-5-yl)oxy]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R, 4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101041-24-0 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[(1H-indol-5-yloxy)methyl]phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

Absolute stereochemistry.

1101041-25-1 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[[(1-methyl-lH-indol-5-yl)oxy]methyl]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101041-29-5 HCAPLUS
3-Furancarboxamide, 4-[[[4-[(1-ethyl-1H-indol-5-yl)methoxy]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101041-30-8 HCAPLUS INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

1101041-31-9 HCAPLUS
3-Furancarboxamide, 4-[[[4-[2-(1H-benzimidazol-l-y1)ethyl]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)- (CA

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101041-32-0 HCAPLUS INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

1101041-33-1 HCAPLUS
3-Furancarboxamide, 4-[[[4-[(2,5-dimethyl-4-thiazolyl)methoxy]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)-(CA INDEX NAME)

Absolute stereochemistry.

1101041-34-2 HCAPLUS
3-Furancarboxamide, 4-[[[4-[2-(1H-benzotriazol-1-y1)ethyl]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)- (CA INDEX

NAME)

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) 1101041-37-5 HCAPLUS 3-Furancatoxamide, tetrahydro-N-hydroxy-4-[[[4-[[2-(1-methylethyl)-4-thiazolyl]methoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

 $1101041-38-6 \quad HCAPLUS \\ 3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[[2-(1-methylethyl)-5-thiazolyl]methoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)$ 

Absolute stereochemistry.

RN CN

1101041-39-7 HCAPLUS
3-Furancarboxanide, 4-[[[4-[(3,5-dimethyl-4-isoxazolyl)methoxy]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-,

(3R,4S)-(CA INDEX NAME)

Absolute stereochemistry.

1101041-40-0 HCAPLUS
3-Furancarboxamide, 4-[[4-(2-butyn-1-yloxy)phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)- (CA INDEX NAME)

02/22/2011

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

Absolute stereochemistry.

1101041-35-3 HCAPLUS
3-Furancarboxamide, 4-[[[4-[(4,5-dimethyl-2-thiazolyl)methoxy]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)-(CA INDEX NAME)

1101041-36-4 HCAPLUS
3-Furancarboxamide, 4-[[[4-[(2,4-dimethyl-5-thiazolyl)methoxy]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)-(CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

Absolute stereochemistry.

$$\operatorname{HO}^{\operatorname{H}} \circ \operatorname{\operatorname{Id}}^{\operatorname{O}} \circ \operatorname{Id}^{\operatorname{O}} \circ \operatorname{\operatorname{Id}}^{\operatorname{O}} \circ \operatorname{\operatorname{Id}}^{\operatorname{O}} \circ \operatorname{Id}^{\operatorname{O}} \circ \operatorname{\operatorname{Id}}^{\operatorname{O}} \circ \operatorname{\operatorname{Id}}^{\operatorname{O}} \circ \operatorname{\operatorname{Id}}^{\operatorname{O}} \circ \operatorname{Id}^{\operatorname{O}} \circ \operatorname{Id}^{\operatorname{O}}$$

1101041-41-1 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(2-pentyn-1-yloxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101041-84-2 HCAPLUS 3-Furancarboxamide, tetrahydro-N-hydroxy-4-[(phenylsulfonyl)methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101041-85-3 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[(4-methylphenyl)sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101041-86-4 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[(4-methoxyphenyl)sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101041-87-5 HCAPLUS
3-Furancarboxamide, 4-[[(4-ethoxyphenyl)sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101041-88-6 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[(4-propoxyphenyl)sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101041-92-2 HCAPLUS
3-Furancarboxamide, 4-[[4-(cyclopropyloxy)phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)-(CA INDEX NAME)

CN

1101041-93-3 HCAPLUS
3-Furancarboxamide, 4-[[[4(cyclobutyloxy)phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)-(CA

INDEX NAME)

Absolute stereochemistry.

1101041-94-4 HCAPLUS
3-Furancarboxamide, 4-[[[4-(cyclopentyloxy)phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)-(CA INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101041-89-7 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(1-methylethoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

1101041-90-0 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(2-methylpropoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

 $\label{local-problem} $$101041-91-1$$ RCAPLUS $$3-Furancarboxamide, $4-[[4-(1,1-dimethylethoxy)phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, $(3R,4S)-dimethylethoxy)phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, $(3R,4S)-dimethylethoxy)phenyl]sulfonyl]sulfonyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, $(3R,4S)-dimethyll]sulfonyl]su$ (CA

INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101041-95-5 HCAPLUS
3-Furancarboxamide, 4-[[[4-(cyclohexyloxy)phenyl]putfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)-

Absolute stereochemistry.

1101041-96-6 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[(4-phenoxyphenyl)sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101041-97-7 HCAPLUS
3-Furancarboxamide, 4-[[[4-(3,5-dimethyl]phenoxy)phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)-(CA INDEX NAME)

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

 $\label{local-problem} $$101041-98-8$$ $$ HCAPLUS$$ 3-Furancarboxamide, $4-[[[4-(3,5-dichlorephenoxy)phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, $(3R,4S)-(CA INDEX NAME)$$$ 

1101041-99-9 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(3-methyl]henoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101042-00-5 HCAPLUS
3-Furancarboxamide, 4-[[4-(3-chlorophenoxy)phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101042-04-9 HCAPLUS
3-Furancarboxamide, 4-[[[4-[(2,6-dimethyl-4-pyridinyl)oxy]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)- (CA INDEX NAME)

1101042-05-0 HCAPLUS INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

1101042-06-1 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(5-quinolinyloxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101042-01-6 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(4-methylphenoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101042-02-7 HCAPLUS
3-Furancarboxamide, 4-[[[4-(4-chlorophenoxy)phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101042-03-8 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(4-pyridinyloxy)phenyl]sulfonyl]methyl]-, (3R, 4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101042-07-2 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(6-quinolinyloxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101042-08-3 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(5-isoquinolinyloxy)phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

Absolute stereochemistry.

1101042-09-4 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)oxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

1101042-10-7 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(phenylmethoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

 $\label{local-equation} $$101042-11-8$$ $$BCAPLUS$$ 3-Furancarboxamide, $4-[[4-[(3,5-dimethylphenyl)methoy]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, $$(3R,4S)-$$$ (CA INDEX NAME)$$$ 

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101042-15-2 HCAPLUS INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

1101042-16-3 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(2-pyridinylmethoxy)phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

Absolute stereochemistry.

1101042-17-4 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(3-pyridinylmethoxy)phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101042-12-9 BCAPLUS
3-Furancarboxamide, 4-[[4-[(3,5-dichlorophenyl)methoxy]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101042-13-0 HCAPLUS
3-Furancarboxamide, 4-[[[4-[(3,5-dimethoxyphenyl)methoxy]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101042-14-1 HCAPLUS
3-Furancarboxamide, 4-[[[4-[(3,5-dibromophenyl)methoxy]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

Absolute stereochemistry.

1101042-18-5 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(4-pyridinylmethoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

1101042-19-6 HCAPLUS
3-Furancarboxamide, 4-[[[4-[(2,6-dimethyl-4-pyridinyl)methoxy]]henyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)-rel- (CA INDEX NAME)

Relative stereochemistry.

RN 1101042-20-9 HCAPLUS

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Page 25

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)
3-Furancarboxamide, 4-[[[4-[(2-chloro-6-methyl-4pyridinyl)methoxy]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)(CA INDEX NAME)

Absolute stereochemistry.

1101042-21-0 HCAPLUS
3-Furancarboxamide, 4-[[[4-[(2-chloro-6-methoxy-4-pyridinyl)methoxy]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)-(CA INDEX NAME)

Absolute stereochemistry.

1101042-22-1 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(4-quinolinylmethoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101042-24-3 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(6-quinolinylmethoxy)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101042-25-4 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101042-26-5 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[(2-methoxy-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R, 48)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101042-23-2 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(5-quinolinylmethoxy)phenyl]sulfonyl]methyl]-, (3K,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101042-27-6 HCAPLUS INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

1101042-28-7 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[2-(4-quinolinyl)ethoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN Absolute stereochemistry. (Continued)

1101042-29-8 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[2-(5-quinolinyl)ethoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

1101042-30-1 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[2-(6-quinoliny1)ethoxy]pheny1]sulfony1]methy1]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101042-31-2 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[2-(5-isoquinolinyl)ethoxy]phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

Absolute stereochemistry.

1101042-32-3 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[2-(2-methyl-4-quinolinyl)ethoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101042-33-4 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[2-(2-methoxy-4-quinolinyl)ethoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

RN 1101042-34-5 HCAPLUS CN INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101042-35-6 HCAPLUS
3-FURANCATROXAMIDe, 4-[[[4-[(3,5-dimethylphenoxy)methyl]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101042-36-7 HCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-(phenoxymethyl)phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) 1101042-37-8 HCAPLUS 3-Furancarboxamide, 4-[[[4-[(3,5-dichlorophenoxy)methyl]phenyl]sulfonyl]methyl]tetrahydro-N-hydroxy-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

1101042-38-9 HCAPLUS INDEX NAME NOT YET ASSIGNED

1101042-39-0 BCAPLUS
3-Furancarboxamide, tetrahydro-N-hydroxy-4-[[[4-[(4-pyridinyloxy)methyl]phenyl]sulfonyl]methyl]-, (3R,48)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

1101042-40-3 HCAPLUS INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

1101043-60-0 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-1-phenyl-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A

1101043-61-1 HCAPLUS CN

3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-1-(2-pyridinyl)-, (3R,4S)-

INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 2-A

PAGE 1-A



1101043-62-2 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinoliny1)methoxy]pheny1]sulfony1]methy1]-1-(3-pyridiny1)-, (3R,4S)-

INDEX NAME)

PAGE 1-A

PAGE 2-A

1101043-63-3 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-1-(4-pyridinyl)-, (3R,4S)-

INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN

(Continued)

PAGE 1-A

PAGE 2-A



1101043-64-4 HCAPLUS
3-Pyrrolidinecarboxamide, 1-acetyl-N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A



1101043-65-5 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinoliny1)methoxy]phenyl]sulfonyl]methyl]-1-(1-oxopropyl)-, (3R,48)-

INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A



1101043-66-6 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-1-(1-oxobutyl)-, (3R,4S)- (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

1101043-67-7 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-1-(2-methyl-1-oxopropyl)-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A

1101043-69-9 HCAPLUS
3-Byrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-1-(2-pyridinylcarbonyl)-,
(3R,4S)- (CR INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A



RN 1101043-68-8 HCAPLUS
CN 3-Pyrrolidinecarboxamide, 1-(2,2-dimethyl-1-oxopropyl)-N-hydroxy-4-[[4[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA
INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A

RN 1101043-70-2 HCAPLUS
CN 3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4quinolinyl)methoxy]phenyl]sulfonyl]methyl]-1-(3-pyridinylcarbonyl)-,
(3R,4S)- (CA INDEX NAME)

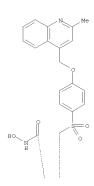
PAGE 1-A

PAGE 2-A

1101043-71-3 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-1-(4-pyridinylcarbonyl)-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)



PAGE 2-A

PAGE 1-A

1101043-73-5 HCAPLUS 3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinoliny])methoxy]phenyl]sulfonyl]methyl]-1-(phenylsulfonyl)-, (3R,4S)-(CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A



1101043-72-4 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-1-(methylsulfonyl)-, (3R,4S)-(CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 2-A

PAGE 1-A

RN 1101043-74-6 BCAPLUS
CN 3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-1-(2-pyridinylsulfonyl)-,
(3R,4S)- (CA INDEX NAME)

PAGE 1-A

PAGE 1-A

PAGE 2-A

1101043-75-7 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinoliny])methoxy]phenyl]phenyl]sulfonyl]-1-(3-pyridinylsulfonyl)-, (3R,43)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 2-A

RN 1101043-77-9 HCAPLUS CN INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A

1101043-76-8 HCAPLUS
3-Fyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinoliny1)methoxy]phenyl]sulfonyl]methyl]-1-(4-pyridinylsulfonyl)-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 2-A

PAGE 1-A

RN 1101043-78-0 HCAPLUS
CN 1-Pyrrolidinecarboxylic acid,
3[(hydroxymaino)carbonyl]-4-[[[4-[(2-methyl4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, propyl ester, (3R,4S)-

INDEX NAME)

PAGE 1-A

PAGE 2-A

RN 1101043-79-1 HCAPLUS
CN 1-Pyrrolidinecarboxylic acid,
3-(Nydroxyamino)carbonyl]-4-[[4-[(2-methyl4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, 1-methylethyl ester,
(3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN

(Continued)

PAGE 1-A

PAGE 2-A

RN 1101043-80-4 HCAPLUS CN INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A

RN 1101043-81-5 HCAPLUS CN INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 2-A

PAGE 1-A

RN 1101043-82-6 HCAPLUS CN INDEX NAME NOT YET ASSIGNED

PAGE 1-A

PAGE 2-A

RN 1101043-83-7 HCAPLUS CN INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A

RN 1101043-84-8 HCAPLUS CN INDEX NAME NOT YET ASSIGNED

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A

1101044-58-9 HCAPLUS
3-Pyrrolidinecarboxamide, 1-ethyl-N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R, 4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 2-A

PAGE 1-A



RN 1101044-59-0 HCAPLUS
CN 3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-1-propyl-, (3R,4S)- (CA INDEX NAME)

PAGE 1-A

PAGE 2-A



1101044-60-3 HCAPLUS
3-Pyrrolidinecarboxamide, 1-butyl-N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R, 4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A



RN 1101044-61-4 HCAPLUS
CN 3-Pyrrolidinecarboxamide,
N-hydroxy-1(-2-methylpropyl)-4-[[[4-[(2-methyl-4quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A

1101044-62-5 HCAPLUS
3-Pyrrolidinecarboxamide, 1-cyclobutyl-N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A



RN 1101044-63-6 HCAPLUS
CN 3-Pyrrolidinecarboxamide, 1-cyclohexyl-N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-, (3R,4S)- (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

1101044-64-7 HCAPLUS 3-Pyrrolidinecarboxamide, 1-cyclopenty1-N-hydroxy-4-[[[4-[(2-methy1-4-quinoliny1)methoxy]pheny1]sulfony1]methy1]-, (3R,4S)- (CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 1-A

PAGE 2-A

OS.CITING REF COUNT: RECORD

THERE ARE 1 CAPLUS RECORDS THAT CITE THIS

REFERENCE COUNT:

FORMAT

(1 CITINGS)
THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE

L5 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

PAGE 2-A

PAGE 1-A

1101044-65-8 HCAPLUS
3-Pyrrolidinecarboxamide, N-hydroxy-4-[[[4-[(2-methyl-4-quinolinyl)methoxy]phenyl]sulfonyl]methyl]-1-(2-propen-1-yl)-, (3R,4S)-(CA INDEX NAME)

Absolute stereochemistry.

L5 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN
ACCESSION NUMBER: 201:876610 HCAPLUS
DOCUMENT NUMBER: 136:19953
ITILE: Preparation of alkynyl aryl sulfonamide hydroxamic acids as TNF-a converting enzyme inhibitors.

Levin, Jeremy I.; Chen, James M.; Zask, Arie
American Cyanamid Company, USA
U.S., 21 pp.
CODEN: USXXAM
DOCUMENT TYPE: CODEN: USXXAM
DAMEDIAN ACCOUNT: English
FAMILY ACC. NUM. COUNT: 1
FAMILY ACC. NUM. COUNT: 1
FATENT INFORMATION:

DOCUMENT TYPE: LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. В1 US 6326516 20011204 US 2000-492980

PRIORITY APPLN. INFO.: US 1999-155250P P 19990127

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT OTHER SOURCE(S): MARPAT 136:19953

AB R8C.tplbond.CCR6R7ZYXNR5CRB1CHR12CONHOH, R8C.tplbond.CCR6R7ZYXNR5CRB11CHR12CONHOH [X = SO2, P(O)R10; Y =

heteroaryl,

REC. TPIROMA. CCREA/EXTRESCRIFICENTECONROH [A = 802, P[O]RLD; Y = Ph. naphthyl; Z = O, NH, CH2, S; R5 = H, alkyl; R6, R7 = H, Me; R8 = H, alkyl, alkenyl, alkynyl, cycloalkyl, heteroaryl, Ph. etc.; R10 = alkyl, cycloalkyl, Ph, heteroaryl; R11, R12 = H, alkyl, cycloalkyl, heteroaryl; Ph. RilR12 = atoms to form (fused) (unsatd.) ring; with provisos], were prepared Thus, [R,28)=2-[[[4-(2-butynyloxy)phenyl]sulfonyl](methyl)amino]-N-hydroxycyclopentanecarboxamide (general preparation given) inhibited TNP-a converting enzyme (TACE) with IC50 = 14 nM.

IT 376630-57-8P, (1R,28)-2-[[[4-(2-Butynyloxy)phenyl]amino]-N-hydroxycyclopentanecarboxamide RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BICL (Biological study); PREP (Preparation); USES (Uses)

(Uses)
(preparation of alkynyl aryl sulfonamide hydroxamic acids as TNF-aconverting enzyme inhibitors)
RN 376630-57-8 HCAPLUS
CN Cyclopentanecarboxamide,
2-[[[4-(2-butyn-1-ylaxy)phenyl]sulfonyl]amino]-N-hydroxy-, (1R,2S)- (CA INDEX NAME)

Absolute stereochemistry.

7 THERE ARE 7 CAPLUS RECORDS THAT CITE THIS

L5 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) (7 CITINGS)
73 THERE ARE 73 CITED REFERENCES AVAILABLE FOR REFERENCE COUNT:

RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

INVENTOR(S): PATENT ASSIGNEE(S): SOURCE:

related compounds as TNF-a converting enzyme (TACE) inhibitors.
Levin, Jeremy Ian; Chen, James Ming; Zask, Arie American Cyanamid Company, USA PCT Int. Appl., 58 pp.
CODEN: PIXXD2
Fatent English 1

2000:535104 HCAPLUS 133:150361

alkynyloxyphenylsulfonylaminoalkylhydroxamic acids

(Continued)

Preparation of

L5 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN ACCESSION NUMBER: 2000:535104 HCAPLUS

DOCUMENT TYPE: LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

DOCUMENT NUMBER:

TITLE:

and

PAT	TENT I	NO.					DATE				ICAT						
WO	2000	0447	11													0000	
		CZ, IN, MD, SK,	DE, IS, MG, SL,	DK, JP, MK, TJ,	DM, KE, MN, TM,	EE, KG, MW, TR,	AZ, ES, KP, MX, TT,	FI, KR, NO, TZ,	GB, KZ, NZ, UA,	GD, LC, PL, UG,	GE, LK, PT, UZ,	GH, LR, RO, VN,	GM, LS, RU, YU,	HR, LT, SD, ZA,	HU, LU, SE, ZW	ID, LV, SG,	IL, MA, SI,
	KW:	DK,	ES,	FI,	FR, GA,	GB,	SD, GR, GW,	IE,	IT,	LU,	MC, SN,	NL, TD,	PT, TG	SE,	BF,	ВJ,	CF,
CA	2356	345			A1		2000	0803		CA 2	000-	2356	345		2	0000	127
EP	1147	078			A1		2001	1024		EP 2	000-	9045	70		2	127	
	R:						ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,
BR	2000							1113		BR 2	000-	7754			2	0000	127
HU	2002	0006	05		A2		2002	0729		HU 2	002-	605			2	0000	127
HU JP	2002	0006 5353	05 83		A3 T		2005 2002			JP 2	000-	5959	68		2	0000	127
NZ	5120	25			A		2003	0829		NZ 2	000-	5120	25		2	0000	127
AU	7694	10			В2		2004	0129		AU 2	-000	2630	6		2	0000	127
ZA	2001	0045	08		A		2002	0902		ZA 2	001-	4508			2	0010	531
NO	2001	0036	39		A		2001	0724		NO 2	001-	3639			2	0010	724
MX	2001	0074	65		A		2001	1203	1	MX 2	001-	7465			2	0010	724
	APP:	LN.	INFO	. :						US 1	999-	2390	83		A 1	9990	127
										WO 2	-000	US18	65		W 2	0000	127

L5 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN

L5 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN OTHER SOURCE(S): MARPAT 133:150361 (Continued)

Title compds. (I; X = SO2, POR10; Y = 5-10 membered heteroaryl, Ph, naphthyl; Z = O, NH, CH2, S; R5 = H, alkyl; R6, R7 = H, Me; R8 = H, alkyl,

alkenyl, alkynyl, cycloalkyl, heteroaryl, heterocycloalkyl, Ph; R9 = H, alkyl, cycloalkyl, Ph; R10 = alkyl, cycloalkyl, Ph, heteroaryl; R11, R12

H, alkyl, cycloalkyl, heteroaryl, heterocycloalkyl; R11R12 = atoms to form

5-10 membered mono- or bicyclic (heterocyclic) ring, Ph, naphthyl; dotted line = optional double bond), were prepared Thus, (IR, 2S) -2-[[4-(2-butynyloxy))phenyl]sulfonyl]methylamino]-N-hydroxycyclopentanecarboxamide [preparation from cis-2-amino-1-cyclopentanecarboxylic acid and 4-(2-butynyloxy)phenylsulfonyl chloride given] inhibited TACE with IC50 = 14 nM.

14 nM. 287096-61-1P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological

(Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of alkynyloxyphenylsulfonylaminoalkylhydroxamic acids and related compds. as TNF-\alpha converting enzyme inhibitors)
RN 287096-61-1 HCAPLUS
CN Cyclopentanecarboxamide,
2-[[[4-(2-butyn-1-yloxy)phenyl]sulfonyl]amino]-N-hydroxy-, (1R,2S)-rel- (CA INDEX NAME)

Relative stereochemistry.

OS.CITING REF COUNT: RECORD

THERE ARE 4 CAPLUS RECORDS THAT CITE THIS

REFERENCE COUNT:

(4 CITINGS) THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE

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02/22/2011

Page 37

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L4 ANSWER 1 OF 5 HCAPLUS COPYRIGHT 2011 ACS on STN ACCESSION NUMBER: 2008:1088521 HCAPLUS

DOCUMENT NUMBER: 149:513677

143:3136// Development of a Suitable Process for the Preparation of a TNF- $\alpha$  Converting Enzyme Inhibitor, WAY-281418 TITLE:

WAY-281418
Wang, Youchu; Papamichelakis, Maria; Chew, Warren;
Sellstedt, John; Noureldin, Razzak; Tadayon, Sam;
Daigmeault, Sylvain
Chemical Development, Wyeth Research, Saint-Laurent,
QC, H4R 176, Can.
Organic Process Research & Development (2008), 12(6),
1253-1260
CODEN: OPRDFK; ISSN: 1083-6160
American Chemical Society
Journal
English
CASREACT 149:513677 AUTHOR(S):

CORPORATE SOURCE:

SOURCE.

PUBLISHER:
DOCUMENT TYPE:
LANGUAGE:
OTHER SOURCE(S):
GI

A suitable process for the preparation of kilogram quantities of a TNF-  $\alpha$  converting enzyme (TACE) inhibitor (WAY-281418) was developed using AB isatin

as starting material and an efficient coupling step for the formation of sulfonamide I in a 15% overall yield. Process preparation of (+)-(1S,2R)-2-aminocyclopentane-1-carboxylic acid (II, (+)-cispentacin),

chiral component for WAY-281418, was successfully scaled up via an asym. hydrogenation reaction. Crystallization allowed the isolation of all intermediates and the final product III.

L4 ANSWER 2 OF 5 HCAPLUS COPYRIGHT 2011 ACS on STN ACCESSION NUMBER: 2006:979177 HCAPLUS DOCUMENT NUMBER: 145:356664 
TITLE: 170 Februarion of  $\beta$ -sulfonamide hydroxamic acid inhibitors of TACE/matrix metalloproteinase Levin, Jeremy I.; Li, Zhong, Diamantidis, George; Lovering, Frank E.; Wang, Weiheng; Condon, Jeffrey S.:

Lin, Yang-I.; Skotnicki, Jerauld S.; Park, Kaapjoo Wyeth, John, and Brother Ltd., USA U.S. Pat. Appl. Publ., 61pp.
CODEN: USXXCO
Patent
English
1 PATENT ASSIGNEE(S):

DOCUMENT TYPE: LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. US 20060211730 US 7595327 PRIORITY APPLN. INFO.: 20060921 20090929 A1 US 2006-377886 US 2005-663785P

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT OTHER SOURCE(S): CASREACT 145:356664; MARPAT 145:356664

AB This invention provides compds. of formula I (wherein J=(un)substituted a monocyclic or bicyclic 5-8 membered cycloalkyl or heterocycloalkyl; R2

a monocyclic or bicyclic 5-8 membered cycloalkyl or heterocycloalkyl; R2 =

H, (un)substituted C1-C6 alkyl, C2-C6 alkenyl or C2-C6 alkynyl; R3 =
(un)substituted naphthyl or bicyclic heteroaryl; R4 and R5 =
independently

H, (un)substituted C1-C6 alkyl, C2-C6 alkenyl or C2-C6 alkynyl; R8 and R9
= independently H, OH, substituted amino, halo, C1-C6 alkyl, etc.; E =
-C1C-, C1N-, N1C-, S or O; X = O, S(O)n, or substituted amino; n = 0-2)
that are useful in treating diseases or disorders mediated by TNF-c, such as arthritis (rheumatoid arthritis (RA) juvenile RA, psoriatic
arthritis, osteoarthritis etc.), tumor metastasis, tissue ulceration,
abnormal wound healing, periodontal disease, bone disease, diabetes
(insulin resistance) and HIV infection, ankylosing spondylitis,
psoriasis,
sepsis, multiple sclerosis, Crohn's disease, degenerative cartilage loss,
asthma, idiopathic pulmonary fibrosis, vasculitis, systemic lupus
erythematosus, irritable bowel syndrome, acute coronary syndrome,
hepatitis C, cachexia, COPD, stroke or type 2 diabetes, and for
alleviation of symptoms thereof. The invention further provides methods
for use of the compds. Preparation of I is exemplified. For example,

L4 ANSWER 1 OF 5 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)
REFERENCE COUNT: 17 THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS

FORMAT

RECORD. ALL CITATIONS AVAILABLE IN THE RE

ANSWER 2 OF 5 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) prepd. by reacting 3-endo-aminobicyclo[2.2.1]hept-5-ene-2-endo-carboxylic acid with 4-(2-methylquinolin-4-ylmethoxy)benzenesulfonyl chloride hydrochloride and reacting the intermediate formed with hydroxylamine.

an assay involving cleavage of pro-TNF by TACE, II had an IC50 of 1.2 nM. OS.CITING REF COUNT: 2 THERE ARE 2 CAPLUS RECORDS THAT CITE THIS

(2 CITINGS)

L4 ANSWER 3 OF 5 HCAPLUS COPYRIGHT 2011 ACS on STN ACCESSION NUMBER: 2002:539654 HCAPLUS DOCUMENT NUMBER:

TITLE:

137:93692
Preparation of
(quinolinylmethoxyphenylsulfonylmethyl)-substituted
pyrrolidinecarboxamides and piperidinecarboxamides as
MMP, TNP, and/or aggreenase inhibitors
Xue, Chu-Biao; Decicco, Carl P.; Re, Xiaohua
Bristol-Myers Squibbl Company Patent Department, USA
PCT Int. Appl., 133 pp.
CODEN: PIXXD2
Patent

WO 2002-US760

W 20020109

INVENTOR (S): PATENT ASSIGNEE(S):

DOCUMENT TYPE: LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION: English

PAT	PATENT NO.  WO 2002055491				KIN	D	DATE			APPL:	ICAT	ION I	NO.		D	ATE	
						-									-		
WO	2002	05549	91		A2		2002	0718		WO 21	002-	US 76	0		21	0020	109
WO	2002	05549	91		A3		2003	0123									
	W:	AE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BY,	BZ,	CA,	CH,	CN,
		co,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,
		GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KZ,	LC,	LK,	LR,
		LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NZ,	OM,	PH,
		PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	TJ,	TM,	TN,	TR,	TT,	TZ,
		UA,	UG,	UZ,	VN,	YU,	ZA,	ZM,	ZW								
	RW:	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AT,	BE,	CH,
		CY,	DE,	DK,	ES,	FI,	FR,	GB,	GR,	IE,	IT,	LU,	MC,	NL,	PT,	SE,	TR,
		BF.	BJ.	CF.	CG,	CI,	CM,	GA.	GN.	GO,	GW,	ML,	MR.	NE.	SN.	TD.	TG
CA	2434	044			A1		2002	0718		CA 21	002-	24341	044		21	0020	109
AU	2002	24698	83		A1		2002	0724		AU 2	002-	2469	83		21	0020	109
US	2003	00878	390		A1		2003	0508		US 21	002-	4354	1		21	0020	109
US	6642	255			B2		2003	1104									
EP	1355	548			A2		2003	1029		EP 2	002-	7147	33		21	0020	109
	R:	AT.	BE.	CH.	DE.	DK.	ES,	FR.	GB,	GR.	IT.	LI.	LU.	NL.	SE.	MC.	PT.
							RO.					-,	- /	-,	-,	-,	-,
PRIORITY	APP				.,	-,	,	,		US 2		2609	57P	1	P 21	0010	111

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT OTHER SOURCE(S): MARPAT 137:93692

L4 ANSWER 3 OF 5 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) exhibited Ki values of ≤ 10 µM against MMP-1, 2, 3, 9, and 13. Thus, I are useful for the treatment of inflammatory disorders and thromboembolic disorder (no data).

OS.CITING REF COUNT: 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS REPORTS.

(1 CITINGS)
THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE REFERENCE COUNT:

FORMAT

L4 ANSWER 3 OF 5 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

Title compds. I [wherein A = COR5, CO2H, CH2CO2H, CO2R6, CONHOH, CONHOR5, CONHOR6, N(OH)CHO, N(OH)COR5, SH, CH2SH, SONHRA, SN2H2Ra, PO3H2, or PO(OH)NHRA; ring B = 3-10 membered (hetero)cycle; Z = absent or (un)substituted (hetero)cycly; U = absent or O, NH, N(alkyl), CO, CO2, CCO, CONH, NHCO, CCO2, etc. X = absent or alkylene, alkenylene, or alkynylene; Y = absent or O, NH, N(alkyl), SOO-2, or CO; Za = (un)substituted (hetero)cyclyl; Rla and Rlb = independently H, alkyl, Ph, PhCH2, CH2OR3, or (un)substituted CH2NH2; or CR1aRlb = (hetero)cyclyl; R2 = Q or (un)substituted alkylene-Q, alkenylene-Q, or alkynylene-Q, Q-substituted alkoxy(alkyl), carbamoyl(alkyl), sulfamoyl(alkyl), etc.;

= H, alkyl, ORa, (un) substituted CH2NH2, or SOO-2Ra; R2b = H or alkyl; Q

H or (un) substituted (hetero) cyclyl; R3 = Q1 or (un) substituted alkylene-Q1, alkenylene-Q1, or alkynylene-Q1, Q1-substituted alkoxy(alky1), carbamoyl(alky1), sulfamoyl(alky1), etc.; or C(R3)2 = (un)substituted (hetcro)cycly1, Q1 = H or (un)substituted Ph, naphthy1,

or

heteroaryl; Ra = H, alkyl, Ph, or PhCH2; p = 0-2; R5 = (un)substituted
alkyl; R6 = phenyl(alkyl), naphthyl, cycloalkyl, alkylcarbonyloxy, etc.;
or pharmaceutically acceptable salt thereof] were prepared as matrix
metalloprotease (MMP), tumor necrosis factor (ThF), and aggrecanase
inhibitors. For example, the
3-(quinolinylmethoxyphenylsulfonylmethyl)-4piperidinecarboxamide (3R,48)-T1-2CF3CO2H was prepared in seventeen
steps starting from the reaction of N-benzyloxycarbonyl-\$\beta\$-alanine and
benzylbronide. Key steps include the cyclization of the 5-aminopentanal
intermediate and the addition of 4-mercaptophenol and
4-chloromethyl-2-methylquinoline\*BCl. \( \lambda\$ number of invention compds.

ANSWER 4 OF 5 HCAPLUS COPYRIGHT 2011 ACS on STN SSION NUMBER: 2001:876610 HCAPLUS

ACCESSION NUMBER: DOCUMENT NUMBER:

UUS COPERIGHT 2011 ACS on STN
2001:876:10 HCAPLUS
136:19953
Preparation of alkynyl aryl sulfonamide hydroxamic
acids as TNT-a converting enzyme inhibitors.
Levin, Jezemy I.; Chen, James M.; Zask, Arie
American Cyanamid Company, USA
U.S., 21 pp.
CODEN: USXXAM
Patent
English
1 INVENTOR(S):

PATENT ASSIGNEE(S): SOURCE:

DOCUMENT TYPE:

LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PATENT NO. KIND US 6326516 PRIORITY APPLN. INFO.: В1 US 2000-492980 20000127 US 1999-155250P P 19990127 20011204

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT OTHER SOURCE(S): MARPAT 136:19953

AB R8C.tplbond.CCR6872YXNR5CRB11CRB12CONHOH,
R8C.tplbond.CCR6872YXNR5CRB11:CR12CONHOH [X = SO2, P(O)R10; Y =

heteroaryl,

roaryl, Ph, naphthyl; Z = O, NH, CH2, S; R5 = H, alkyl; R6, R7 = H, Me; R8 = H, alkyl, alkenyl, alkynyl, cycloalkyl, heteroaryl, Ph, etc.; R10 = alkyl, cycloalkyl, Ph, heteroaryl; R11, R12 = H, alkyl, cycloalkyl, heteroaryl; Ph; R1R12 = atoms to form (fused) (unsatd.) ring; with provisos], were prepared Thus, (1R, 2S)-2-[[[4-(2-butynyloxy)phenyl]sulfonyl](methyl)amino]-

(18,25)-2-[[[4-(2-Dutynyloxy/)pnenyljsulronyl](metnyl/amino)-N-hydroxycyclopentanecarboxamide (general preparation given) inhibited TNF-a converting enzyme (TACE) with IC50 = 14 mM. OS.CITING REF COUNT: 7 THERE ARE 7 CAPIUS RECORDS THAT CITE THIS

RECORD

(7 CITINGS)
THERE ARE 73 CITED REFERENCES AVAILABLE FOR 7.3 REFERENCE COUNT:

RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

L4 ANSWER 5 OF 5 HCAPLUS COPYRIGHT 2011 ACS ON STN ACCESSION NUMBER: 2000:535104 HCAPLUS DOCUMENT NUMBER: 133:150361 TITLE: Preparation of

alkynyloxyphenylsulfonylaminoalkylhydroxamic acids

related compounds as TNF- $\alpha$  converting enzyme (TACE) inhibitors. Levin, Jeremy Ian; Chen, James Ming; Zask, Arie American Cyanamid Company, USA PCT Int. Appl., 58 pp. CODEN: PIXXD2 Patent English 1 INVENTOR(S):
PATENT ASSIGNEE(S):
SOURCE:

DOCUMENT TYPE: LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

	ENT																ATE	
	2000																0000	127
								BA,										
								FI,										
								KR,										
		MD,	MG,	MK,	MN,	MW,	MX,	NO,	NZ,	PI	, P.	, R	ο,	RU,	SD,	SE,	SG,	SI,
		SK,	SL,	TJ,	TM,	TR,	TT,	TZ,	UA,	UG	, U2	, v	N,	YU,	ZA,	ZW		
	RW:	GH,	GM,	KE,	LS,	MW,	SD,	SL,	SZ,	T2	, U	, Zi	N,	AT,	BE,	CH,	CY,	DE,
		DK,	ES,	FI,	FR,	GB,	GR,	IE,	IT,	LU	J, M	, N	L,	PT,	SE,	BF,	ΒJ,	CF,
		CG,	CI,	CM,	GΑ,	GN,	GW,	ML,	MR,	NE	i, Si	I, T	D,	TG				
	2356							0803										
EP	1147																	
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			SI,															
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	2001				A		2002										0010	
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	2001				Α		2001	1203									0010	
(TI	APP	LN.	INFO	. :						US	1999	-23	308	33		A 1	9990	127

WO 2000-US1865

W 20000127

OTHER SOURCE(S): MARPAT 133:150361

L4 ANSWER 5 OF 5 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

AB Title compds. (I; X = SO2, POR10; Y = 5-10 membered heteroaryl, Ph, naphthyl; Z = O, NH, CH2, S; R5 = H, alkyl; R6, R7 = H, Me; R8 = H, alkyl, alkenyl, alkynyl, cycloalkyl, heteroaryl, heterocycloalkyl, Ph; R9 = H, alkyl, cycloalkyl, Ph; R10 = alkyl, cycloalkyl, Ph, heteroaryl; R11, R12

H, alkyl, cycloalkyl, heteroaryl, heterocycloalkyl; R11R12 = atoms to

form
5-10 membered mono- or bicyclic (heterocyclic) ring, Ph, naphthyl; dotted
line = optional double bond), were prepared Thus,
(IR, ZS)-2-[[[4-(2-butynyloxy)phenyl]sulfonyl]methylamino]-Nhydroxycyclopentanecarboxamide [preparation from
cis-2-amino-1-cyclopentanecarboxylic acid and
4-(2-butynyloxy)phenylsulfonyl chloride given] inhibited TACE with IC50 =
14 nM.

OS.CITING REF COUNT: 4 THERE ARE 4 CAPLUS RECORDS THAT CITE THIS
RECORD
(4 CITINGS)

 $(4\ \mbox{CITINGS})$  There are 8 cited references available for this record. All citations available in the re REFERENCE COUNT:

FORMAT

=> FIL REGISTRY

COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 69.76 270.35

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL ENTRY SESSION CA SUBSCRIBER PRICE -6.96 -6.96

FILE 'REGISTRY' ENTERED AT 09:55:08 ON 22 FEB 2011 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2011 American Chemical Society (ACS)

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 21 FEB 2011 HIGHEST RN 1263357-65-8 DICTIONARY FILE UPDATES: 21 FEB 2011 HIGHEST RN 1263357-65-8

New CAS Information Use Policies, enter HELP USAGETERMS for details.

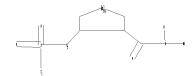
TSCA INFORMATION NOW CURRENT THROUGH June 26, 2010.

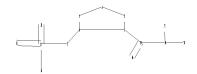
Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

Uploading C:\Program Files\Stnexp\Queries\10593748a.str





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chain nodes :
7 8 11 12 13 14 15 16 17 18
ring nodes :
1 2 3 4 5
chain bonds :
3-14 4-7 7-8 8-11 8-12 8-13 14-15 14-16 15-17 15-18
ring bonds :
1-2 1-5 2-3 3-4 4-5
exact/norm bonds :
4-7 7-8 8-11 8-12 8-13 14-15 14-16 15-18
exact bonds :
1-2 1-5 2-3 3-4 3-14 4-5 15-17
isolated ring systems :
containing 1 :
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G1:0,S,N,CH2

G2:CH2,NH

G3:Cb, Cy, Hy, Ak, Ph

Match level:

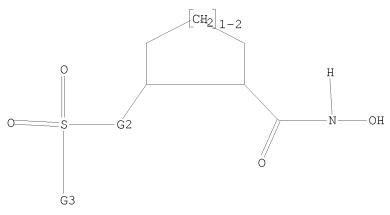
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 7:CLASS 8:CLASS 11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:CLASS 18:CLASS

L6 STRUCTURE UPLOADED

=> d 16

L6 HAS NO ANSWERS

L6



G1 O, S, N, CH2

G2 CH2, NH

G3 Cb,Cy,Hy,Ak,Ph

Structure attributes must be viewed using STN Express query preparation.

=> s 16

SAMPLE SEARCH INITIATED 09:55:42 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 14 TO ITERATE

14 ITERATIONS 100.0% PROCESSED 6 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\* BATCH \*\*COMPLETE\*\*

PROJECTED ITERATIONS: 56 TO 504

PROJECTED ANSWERS: 6 TO 266

6 SEA SSS SAM L6

=> s 16 sss full

FULL SEARCH INITIATED 09:55:48 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 154 TO ITERATE

36 ANSWERS 100.0% PROCESSED 154 ITERATIONS

SEARCH TIME: 00.00.01

36 SEA SSS FUL L6 L8

=> FIL HCAPLUS

SINCE FILE TOTAL ENTRY SESSION COST IN U.S. DOLLARS FULL ESTIMATED COST 196.86 467.21

DISCOUNT AMOUNTS (FOR OUALIFYING ACCOUNTS) SINCE FILE TOTAL SESSION ENTRY

CA SUBSCRIBER PRICE 0.00 -6.96FILE 'HCAPLUS' ENTERED AT 09:55:52 ON 22 FEB 2011

USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2011 AMERICAN CHEMICAL SOCIETY (ACS)

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FILE COVERS 1907 - 22 Feb 2011 VOL 154 ISS 9 FILE LAST UPDATED: 21 Feb 2011 (20110221/ED) REVISED CLASS FIELDS (/NCL) LAST RELOADED: Oct 2010 USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Oct 2010

HCAplus now includes complete International Patent Classification (IPC) reclassification data for the fourth quarter of 2010.

CAS Information Use Policies apply and are available at:

http://www.cas.org/legal/infopolicy.html

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 18L9 7 L8

=> s 19 and py<=2004 25160617 PY<=2004 3 L9 AND PY<=2004  $T_{1}10$ 

=> d l10 ibib abs hitstr tot

L10 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN ACCESSION NUMBER: 2001:876610 HCAPLUS

DOCUMENT NUMBER: 136:19953 TITLE:

136:19953
Preparation of alkynyl aryl sulfonamide hydroxamic acids as TNF-a converting enzyme inhibitors.
Levin, Jereny I.; Chen, James M.; Zask, Arie American Cyanamid Company, USA
U.S., 21 pp.
CODEN: USXXAM INVENTOR (S)

PATENT ASSIGNEE(S): SOURCE:

DOCUMENT TYPE: English

LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE US 6326516 В1 20011204 US 2000-492980 20000127

PRIORITY APPLN. INFO.: US 1999-155250P P 19990127

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT OTHER SOURCE(S): MARPAT 136:19953
AB 88C.tplbond.CCR6872YXNR5CR11CR12CONHOH,
R8C.tplbond.CCR6872YXNR5CR11;CR12CONHOH [X = SO2, P(O)R10; Y =

heteroaryl,

R8C.tplbond.CCR6R7EXNRSCR11:cR12CONNOM [K = 802, P(O)R10; Y = heteroaryl, Ph, naphtyl; Z = O, NH, CH2, S; R5 = H, alkyl; R6, R7 = H, Me; R8 = H, alkyl, alkenyl, alkynyl, cycloalkyl, heteroaryl, Ph, etc.; R10 = alkyl, cycloalkyl, Ph, heteroaryl; R11, R12 = H, alkyl, cycloalkyl, heteroaryl; Ph; R11R12 = atoms to form (fused) (unsatd.) ring; with provisos], were prepared Thus, [R,28)-2-[[[4-(2-butynyloxy)phenyl]sulfonyl](methyl)amino]N-hydroxycyclopentanecarboxamide (general preparation given) inhibited TNP-m converting enzyme (TACE) with ICSO = 14 nM.

IT 287096-63-3P, (cis)-2-[[[4-(2-Butynyloxy)phenyl]sulfonyl]amino]-N-hydroxycyclohexanecarboxamide 376630-56-TP, (1R,2R)-2-[[[4-(2-Butynyloxy)phenyl]sulfonyl]amino]-N-hydroxycyclohexanecarboxamide 376630-59-0P, (IR,28)-2-[[[4-(2-Butynyloxy)phenyl]sulfonyl]amino]-N-hydroxycyclopentanecarboxamide 376630-59-0P, (IR,2R,3S,4R)-3-([[4-(2-Butynyloyy)phenyl]sulfonyl]amino]-N-hydroxybicyclo[2.2.1]heptan-2-carboxamide RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

20/09-63-3 HCAPLUS
Cyclohexanecarboxamide, 2-[[[4-(2-butyn-1-yloxy)phenyl]sulfonyl]amino]-N-hydroxy-, (1R,2S)-rel- (CA INDEX NAME)

Relative stereochemistry.

L10 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

S.CITING REF COUNT: THERE ARE 7 CAPLUS RECORDS THAT CITE THIS

(7 CITINGS)
THERE ARE 73 CITED REFERENCES AVAILABLE FOR REFERENCE COUNT:

RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

L10 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

376630-56-7 HCAPLUS Cyclohexanecarboxamide, 2-[[[4-(2-butyn-1-yloxy)phenyl]sulfonyl]amino]-N-hydroxy-, (1R,2R)- (CA INDEX NAME)

Absolute stereochemistry.

376630-57-8 HCAPLUS

Cyclopentanecarboxamide, [4-(2-butyn-1-yloxy)phenyl]sulfonyl]amino]-N-hydroxy-, (1R,2S)- (CA INDEX NAME) 2-[[[4-

Absolute stereochemistry.

$$\mathsf{Me-c} = \mathsf{C} \qquad \qquad \mathsf{S} \qquad \mathsf{NH} \qquad \mathsf{OH}$$

376630-59-0 HCAPLUS
Bicyclo[2.2.1]heptane-2-carboxamide,
3-[[[4-(2-butyn-1-yloxy)phenyl]sulfonyl]amino]-N-hydroxy-,

(CA INDEX NAME)

Absolute stereochemistry.

L10 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN ACCESSION NUMBER: 2000:553546 HCAPLUS DOCUMENT NUMBER: 133:150358

2000:553546 HCAPLUS
133:150358
Preparation of sulfonamide derivatives having cyclic structures as matrix metalloprotease inhibitors and TNF production inhibitors
Watanabe, Fuminiko; Tsuzuki, Hiroshige
Shionogi and Co. Ltd., Japan
PCT Int. Appl., 87 pp.
CODEN: PIXXD2
Patent
Japanese
1

INVENTOR (S) .

PATENT ASSIGNEE(S): SOURCE:

DOCUMENT TYPE: LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

	PAT:	ENT :	NO.			KIN	D	DATE		APPLICATION NO.							DATE			
							_									_				
	WO :	2000	0461	89		A1		2000	0810	,	WO 2	000-	JP44	6		2	0000	128		
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		W:	ΑE,	AL,	AM,	AT,	ΑU,	AZ,	BA,	BB,	BG,	BR,	BY,	CA,	CH,	CN,	CR,	CU,		
			CZ,	DE,	DK,	DM,	EE,	ES,	FI,	GB,	GD,	GE,	GH,	GM,	HR,	HU,	ID,	IL,		
		IN, IS, JP					KG,	KR,	KZ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,		
	MG, MK, MN					MW,	MX,	NO,	NZ,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,		
	SL, TJ, TM					TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VN,	YU,	ZA,	ZW				
		RW:	GH,	GM,	KE,	LS,	MW,	SD,	SL,	SZ,	TZ,	UG,	ZW,	AT,	BE,	CH,	CY,	DE,		
			DK,	ES,	FI,	FR,	GB,	GR,	IE,	IT,	LU,	MC,	NL,	PT,	SE,	BF,	ВJ,	CF,		
	CG, CI, CM						GN,	GW,	ML,	MR,	NE,	SN,	TD,	TG						
PRIO	RITY	APP	IN.	TNFO	. :						TP 1	999-	2505		A 19990202					

MARPAT 133:150358 OTHER SOURCE(S):

AB The title derivs. I [A is Q1, etc.; R1 is COOR6 or the like; R6 is hydrogen, alkyl; R2 is optionally substituted arylene or the like; R3 is CONH or the like; R4 is optionally substituted aryl or the like; R5 is hydrogen or the like; and m and n are each independently 0 or 1] are prepared The title compound II in vitro showed IC50 of 9.7 µM against MMP-8. Formulations are given.

IT 287395-23-7P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological

L10 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (USEA) (prepn. of sulfonanide derivs. having cyclic structures as matrix metalloprotease inhibitors and TNF prodn. inhibitors)
RN 287395-23-7 HCAPLUS
CN Bicyclo[2,2.1]hept-5-ene-2-carboxamide, 3-[[5-[4-(dimthylamino)phenyl]-2-thienyl]sulfonyl]amino]-N-hydroxy-, (18,28,38,4R)- (CA INDEX NAME)

Absolute stereochemistry.

S.CITING REF COUNT: THERE ARE 5 CAPLUS RECORDS THAT CITE THIS

(6 CITINGS) THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE REFERENCE COUNT:

FORMAT

ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN R SOURCE(S): MARPAT 133:150361 (Continued) OTHER

Title compds. (I; X = SO2, POR10; Y = 5-10 membered heteroaryl, Ph, naphthyl; Z = O, NH, CH2, S; R5 = H, alkyl; R6, R7 = H, Me; R8 = H, alkyl,

alkenyl, alkynyl, cycloalkyl, heteroaryl, heterocycloalkyl, Ph; R9 = H, alkyl, cycloalkyl, Ph; R10 = alkyl, cycloalkyl, Ph, heteroaryl; R11, R12

H, alkyl, cycloalkyl, heteroaryl, heterocycloalkyl; R11R12 = atoms to form

5-10 membered mono- or bicyclic (heterocyclic) ring, Ph, naphthyl; dotted line = optional double bond), were prepared Thus, (IR, 2S) -2-[[[4-(2-butynyloxy)phenyl]sulfonyl]methylamino]-N-hydroxycyclopentanecarboxamide [preparation from cis-2-amino-1-cyclopentanecarboxylic acid and 4-(2-butynyloxy)phenylsulfonyl chloride given] inhibited TACE with IC50 = 14 nM.

14 nM. 287096-58-6P 287096-65-5P 287096-63-3P 287096-61-1P IT

RL: BAC (Biological activity or effector, except adverse); BSU

(Biological

study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of alkynyloxyphenylsulfonylaminoalkylhydroxamic acids and related compds. as TNP-α converting enzyme inhibitors) 281096-58-6 HCAPLUS (Cyclohexanecarboxamide, 2-[[[4-(2-butyn-1-yloxy)phenyl]sulfonyl]amino]-N-hydroxy-, (IR,2R)-rel- (CA INDEX NAME)

Relative stereochemistry.

RN 287096-61-1 HCAPLUS CN Cyclopentanecarboxamide, 2-[[[4-(2-butyn-1-yloxy)phenyl]sulfonyl]amino]-N-hydroxy-, (1R,2S)-rel- (CA INDEX NAME)

L10 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN ACCESSION NUMBER: 2000:535104 HCAPLUS

DOCUMENT NUMBER: 133:150361

Preparation of TITLE:

alkynyloxyphenylsulfonylaminoalkylhydroxamic acids and

related compounds as TNF- $\alpha$  converting enzyme (TACE) inhibitors. Levin, Jeremy Ian; Chen, James Ming; Zask, Arie American Cyanamid Company, USA PCT Int. Appl., 58 pp. CODEN: PIXXD2 Patent English 1

INVENTOR (S) PATENT ASSIGNEE(S):

SOURCE:

DOCUMENT TYPE: LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

											LICAT							
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											, MC,			SE,	BF,	ВJ,	CF,	
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		_					 											
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2	A	2001	0045	08		A	2002	0902		ZA	2001-	4508			2	0010	531	
_						_												
1	10	2001	0036	39		A	2001	0724		NO	2001-	3639			- 2	0010	724	
N	ſΧ	2001	0074	65		A	2001	1203		MX	2001-	7465			2	0010	724	
RI	TY	APP	LN.	INFO	. :					US	1999-	2390	83		A 1	9990	127	
										WO.	2000-	11518	65	,	w 2	nnnn	127	
											2000-	0010	~~			0000	16/	

L10 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

287096-63-3 HCAPLUS Cyclohexanecarboxamide, 2-[[[4-(2-butyn-1-yloxy)phenyl]sulfonyl]amino]-N-hydroxy-, (IR,28)-rel- (CA INDEX NAME)

287096-65-5 HCAPLUS
Bicyclo[2.2.1]heptane-2-carboxamide,
3-[[4-(2-butyn-1-yloxy)phenyl]sulfonyl]amino]-N-hydroxy-,
(1R,2S,3R,4S)-rel- (CA INDEX NAME)

Relative stereochemistry.

OS.CITING REF COUNT: RECORD THERE ARE 4 CAPLUS RECORDS THAT CITE THIS

(4 CITINGS)
THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE REFERENCE COUNT:

FORMAT

L10 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued)

=> d 19 ibib abs tot

L9 ANSWER 1 OF 7 HCAPLUS COPYRIGHT 2011 ACS on STN ACCESSION NUMBER: 2008:1088521 HCAPLUS

DOCUMENT NUMBER: 149:513677

Parallope // Development of a Suitable Process for the Preparation of a TNF- $\alpha$  Converting Enzyme Inhibitor, WAY-281418 TITLE:

WAY-281418
Wang, Youchu; Papamichelakis, Maria; Chew, Warren;
Sellstedt, John; Noureldin, Razzak; Tadayon, Sam;
Daigmeault, Sylvain
Chemical Development, Wyeth Research, Saint-Laurent,
QC, H4R 176, Can.
Organic Process Research & Development (2008), 12(6),
1253-1260
CODEN: OPRDFK; ISSN: 1083-6160
American Chemical Society
Journal
English
CASREACT 149:513677 AUTHOR(S):

CORPORATE SOURCE.

SOURCE.

PUBLISHER:
DOCUMENT TYPE:
LANGUAGE:
OTHER SOURCE(S):
GI

AB A suitable process for the preparation of kilogram quantities of a TNF-  $\alpha$  converting enzyme (TACE) inhibitor (WAY-281418) was developed using isatin

as starting material and an efficient coupling step for the formation of sulfonamide I in a 15% overall yield. Process preparation of (+)-(18,2R)-2-aminocyclopentane-1-carboxylic acid (II, (+)-cispentacin),

chiral component for WAY-281418, was successfully scaled up via an asym.

L9 ANSWER 2 OF 7 HCAPLUS COPYRIGHT 2011 ACS on STN ACCESSION NUMBER: 2006:979177 HCAPLUS DOCUMENT NUMBER: 145:35664 
TITLE: 6 Februarion of  $\beta$ -sulfonamide hydroxamic acid inhibitors of TACE/matrix metalloproteinase INVENTOR(S): Levin, Jeremy I.; Li, Zhong, Diamantidis, George; Lovering, Frank E.; Wang, Weiheng; Condon, Jeffrey S.:

Lin, Yang-I.; Skotnicki, Jerauld S.; Park, Kaapjoo Wyeth, John, and Brother Ltd., USA U.S. Pat. Appl. Publ., 61pp.
CODEN: USXXCO
Patent
English
1 PATENT ASSIGNEE(S):

SOURCE:

DOCUMENT TYPE: LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

10593748.trn

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 20060211730	A1	20060921	US 2006-377886	20060316
US 7595327	B2	20090929		
PRIORITY APPLN. INFO.:			US 2005-663785P P	20050321

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT OTHER SOURCE(S): CASREACT 145:356664; MARPAT 145:356664 OTHER SOURCE(S):

AB This invention provides compds. of formula I (wherein J=(un)substituted a monocyclic or bicyclic 5-8 membered cycloalkyl or heterocycloalkyl; R2

H, (un)substituted C1-C6 alkyl, C2-C6 alkenyl or C2-C6 alkynyl; R3 = (un)substituted naphthyl or bicyclic heteroaryl; R4 and R5 =

H, (un) substituted capthyl or bicyclic heteroaryl; R4 and R5 = independently

H, (un) substituted c1-C6 alkyl, C2-C6 alkenyl or C2-C6 alkynyl; R8 and R9 = independently H, OH, substituted amino, halo, C1-C6 alkyl, etc.; E = -C1C-, -C1N-, -N1C-, S or O; X = O, S(O)n, or substituted amino; n = 0-2) that are useful in treating diseases or disorders mediated by TNF-α, such as arthritis (rheumatoid arthritis (RA), juvenile RA, psoriatic arthritis, osteoarthritis etc.), tumor metastasis, tissue ulceration, abnormal wound healing, periodontal disease, bone disease, diabetes (insulin resistance) and HTV infection, ankylosing spondylitis, psoriasis, sepsis, multiple sclerosis, Crohn's disease, degenerative cartilage loss, asthma, idiopathic pulmonary fibrosis, vasculitis, systemic lupus erythematosus, irritable bowel syndrome, acute coronary syndrome, hepatitis C, cachexia, COPD, stroke or type 2 diabetes, and for

02/22/2011 Page 50 L9 ANSWER 1 OF 7 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) hydrogenation reaction. Crystn. allowed the isolation of all intermediates and the final product III.

REFERENCE COUNT: 17 THERE ARE 17 CITED REFERENCES AVAILABLE FOR

THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 2 OF 7 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) alleviation of symptoms thereof. The invention further provides methods for use of the compds. Frepn. of I is exemplified. For example, II was prepd. by reacting 3-endo-aminobicyclo[2.2.1]hept-5-ene-2-endo-carboxylic acid with 4-(2-methylquinolin-4-ylmethoxy)benzenesulfonyl chloride hydrochloride and reacting the intermediate formed with hydroxylamine.

an assay involving cleavage of pro-TNF by TACE, II had an IC50 of 1.2 nM. OS.CITING REF COUNT: 2 THERE ARE 2 CAPLUS RECORDS THAT CITE THIS

(2 CITINGS)

FORMAT

L9 ANSWER 3 OF 7 HCAPLUS COPYRIGHT 2011 ACS on STN ACCESSION NUMBER: 2006:757211 HCAPLUS

DOCUMENT NUMBER: 145:241166

145:241166
Targeting ADAM-mediated ligand cleavage to inhibit
HER3 and EGFR pathways in non-small cell lung cancer
Zhou, Bin-Bing S.; Peyton, Michael; He, Biao; Liu,
Changnian; Girard, Luc; Caudler, Elan; Lo, Yvonne;
Baribaud, Frederic; Mikami, Iwao; Requatr, Noemi;
Yang, Gengjie; Li, Yanlong; Yao, Wenqing; Vaddi, TITLE: AUTHOR(S):

Krie. Gazdar, Adi F.; Friedman, Steven M.; Jablons, David M.; Newton, Robert C.; Fridman, Jordan S.; Minna,

John

CORPORATE SOURCE:
Drug Discovery, Experimental Station, Incyte
Corporation, Wilmington, DE, 19880, USA
SOURCE:
Cancer Cell (2006), 10(1), 39-50
COEN: CCAECI; ISSN: 1535-6108
PUBLISHER:
Cell Press
DOCUMENT TYPE:
Journal
LANGUAGE:
English
AB We describe here the existence of a heregulin-HER3 autocrine loop, and the

contribution of heregulin-dependent, HER2-mediated HER3 activation to gefitinib insensitivity in non-small cell lung cancer (NSCLC). ADAM17 protein, a major ErbB ligand sheddase, is upregulated in NSCLC and is required not only for heregulin-dependent HER3 signaling, but also for EGFR ligand-dependent signaling in NSCLC cell lines. A selective ADAM inhibitor, INCB3619, prevents the processing and activation of multiple ErbB ligands, including heregulin. In addition, INCB3619 inhibits egfitinib-resistant HER3 signaling and enhances gefitinib inhibition of EGFR signaling in NSCLC. These results show that ADAM inhibition affects multiple ErbB pathways in NSCLC and thus offers an excellent opportunity for pharmacol. intervention, either alone or in combination with other drugs.

drugs. OS.CITING REF COUNT: 98 THERE ARE 98 CAPLUS RECORDS THAT CITE THIS

RECORD (99 CITINGS)
THERE ARE 60 CITED REFERENCES AVAILABLE FOR 60 REFERENCE COUNT:

RECORD. ALL CITATIONS AVAILABLE IN THE RE

ANSWER 4 OF 7 HCAPLUS COPYRIGHT 2011 ACS on STN

Title compds. I [X = (CH2)nO, (CH2)nS, (CH2)n(CH2), etc.; n = 0-2; R = (un)substituted alkyl, alkenyl, alkynyl, etc.; Z = NH or CH2] and their pharmaceutically acceptable salts, are prepared and disclosed as AB inhibitors

inhibitors

of matrix metalloproteinases (MMP). Thus, e.g., II was prepared by sulfonylation of cis-2-amino-1-cyclohexanecarboxylic acid with biphenyl-4-sulfonyl chloride and subsequent amidation using O-(trimethylsily)lnydroxylamine. The inhibitory activity of I towards MMP-2, MMP-3 and MMP-9 was evaluated using fluorometric substrate-degradation assays and it was revealed that selected compds. of the invention displayed IC50 values in the range of 125 up to 150 nM against MMP-2, 145 up to 175 nM against MMP-9 and above 3000 nM against MMP-3. I as inhibitors of matrix metalloproteinases should prove useful in the treatment of leukemia, melanoma and carcinoma. Pharmaceutical compns. comprising I are disclosed.

OS.CITING REF COUNT: 2 THERE ARE 2 CAPLUS RECORDS THAT CITE THIS RECORD

(2 CITINGS)
THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE REFERENCE COUNT:

FORMAT

L9 ANSWER 4 OF 7 HCAPLUS COPYRIGHT 2011 ACS on STN ACCESSION NUMBER: 2005:1075765 HCAPLUS

DOCUMENT NUMBER: 143:346847

143:346847
Preparation of cyclohexylcarboxamide derivatives as inhibitors of matrix metalloproteinases
Ananthan, Subramaniam
Southern Research Institute, USA
PCT Int. Appl., 52 pp.
CODEN: PIXMD2 TITLE:

INVENTOR(S):

PATENT ASSIGNEE(S): SOURCE:

DOCUMENT TYPE:

LANGUAGE: FAMILY ACC. NUM. COUNT: English

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	WO											2005-						
		W:										, BG,						
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												, SC, US,						
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		RW:	BW.	GH,	GM,	KE.	LS,	MW.	MZ.	NA.	SD	. SL.	SZ.	TZ.	UG,	ZM.	ZW.	AM.
			AZ,	BY,	KG,	KZ,	MD,	RU,	TJ,	TM,	AT.	, BE,	BG,	CH,	CY,	CZ,	DE,	DK,
			EE,	ES,	FI,	FR,	GB,	GR,	HU,	IE,	IS,	, IT,	LT,	LU,	MC,	NL,	PL,	PT,
			RO,	SE,	SI,	SK,	TR,	BF,	ВJ,	CF,	CG,	, CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,
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	EP	1735						2009										
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	US	2008	0312	329		A1		2008	1218			2008-						
PRIC	ORIT:	APP	LN.	INFO	. :						US :	2004-	5553	80P		P 2	0040	322
											WO:	2005-	0892	63		w 2	UU 50	321

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT
OTHER SOURCE(S): CASREACT 143:346847; MARPAT 143:346847 OTHER SOURCE(S):

L9 ANSWER 5 OF 7 HCAPLUS COPYRIGHT 2011 ACS ON STN ACCESSION NUMBER: 2001:876610 HCAPLUS DOCUMENT NUMBER: 136:19953

LUS COPYRIGHT 2011 ACS on STN
2001:876610 HCAPLUS
136:19953
Preparation of alkynyl aryl sulfonamide hydroxamic
acids as TNR-a converting enzyme inhibitors.
Levin, Jeremy I.; Chen, James M.; Zask, Arie
American Cyanamid Company, USA
U.S., 21 pp.
CODEN: USXXAM
Patent
English
1

INVENTOR(S):

PATENT ASSIGNEE(S):

DOCUMENT TYPE:

LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6326516	B1	20011204	US 2000-492980	20000127
PRIORITY APPIN. INFO.:			HS 1999-155250P P	19990127

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT OTHER SOURCE(S): MARPAT 136:19953

AB R8C.tplbond.CCR6872YXNR5CRB11CRB12CONHOH,
R8C.tplbond.CCR6872YXNR5CRB11:CR12CONHOH [X = SO2, P(O)R10; Y =

heteroaryl,

coaryl, Ph, naphthyl; Z = O, NH, CH2, S; R5 = H, alkyl; R6, R7 = H, Me; R8 = H, alkyl, alkenyl, alkynyl, cycloalkyl, heteroaryl, Ph, etc.; R10 = alkyl, cycloalkyl, Ph, heteroaryl; R11, R12 = H, alkyl, cycloalkyl, heteroaryl, Ph; R11R12 = atoms to form (fused) (unsatd.) ring; with provisos], were prepared Thus, (1R, 2S)-2-[[[4-(2-butynyloxy)phenyl]sulfonyl](methyl)amino]-

(18,25)-2-[[[4-(2-Dutynyloxy/)pnenyljsulronyl](metnyl/amino)-N-hydroxycyclopentanecarboxamide (general preparation given) inhibited TNF-a converting enzyme (TACE) with IC50 = 14 mM. OS.CITING REF COUNT: 7 THERE ARE 7 CAPIUS RECORDS THAT CITE THIS

RECORD

(7 CITINGS)
THERE ARE 73 CITED REFERENCES AVAILABLE FOR 73 REFERENCE COUNT:

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

(Continued)

L9 ANSWER 6 OF 7 HCAPLUS COPYRIGHT 2011 ACS on STN ACCESSION NUMBER: 2000:553546 HCAPLUS 133:150358
Preparation of sulfonamide derivatives having cyclic structures as matrix metalloprotease inhibitors and TNF production inhibitors watanabe, Fumihiko; Tsuzuki, Hiroshige Shionogi and Co., Ltd., Japan PCT Int. Appl., 87 pp. CODEN: PIXD2
Patent
Japanese DOCUMENT NUMBER: 133:150358 TITLE: INVENTOR(S): PATENT ASSIGNEE(S): SOURCE: DOCUMENT TYPE: LANGHAGE . LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE W1 2000046189 A1 20000810 W2 2000-JP446 20000128
W1 AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU,
CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL,
IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, AA, MD,
MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK,
SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, VU, ZA, ZW
RN: GH, CM, KE, LS, MN, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
PRIORITY APPLN: INFO:

OTHER SOURCE(S): MARPAT 133:150358

(CH<sub>2</sub>)<sub>m</sub>-R<sup>1</sup> (CH<sub>2</sub>)<sub>n</sub>-N-SO<sub>2</sub>-R<sup>2</sup>-R<sup>3</sup>-R<sup>4</sup> CO2H NH-SO2

AB The title derivs. I [A is Q1, etc.; R1 is COOR6 or the like; R6 is hydrogen, alkyl; R2 is optionally substituted arylene or the like; R3 is CONH or the like; R4 is optionally substituted aryl or the like; R5 is hydrogen or the like; and m and n are each independently 0 or 1] are prepared The title compound II in vitro showed IC50 of 9.7 µM against MMP-8. Formulations are given.

S.CITING REF COUNT: 5 THERE ARE 5 CAPLUS RECORDS THAT CITE THIS RECORD

L9 ANSWER 7 OF 7 HCAPLUS COPYRIGHT 2011 ACS on STN
ACCESSION NUMBER: 2000:535104 HCAPLUS
DOCUMENT NUMBER: 133:150361
TITLE: Preparation of alkynyloxyphenylsulfonylaminoalkylhydroxamic acids

related compounds as TNF- $\alpha$  converting enzyme (TACE) inhibitors. Levin, Jeremy Ian; Chen, James Ming; Zask, Arie American Cyanamid Company, USA PCT Int. Appl., 58 pp. CODEN: PIXXD2 Patent English 1 INVENTOR (S) . PATENT ASSIGNEE(S): SOURCE:

DOCUMENT TYPE: LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

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		2001				A		2001	1203			2001					0010	
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OTHER SOURCE(S): MARPAT 133:150361

L9 ANSWER 6 OF 7 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) (6 CITINGS)
THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE REFERENCE COUNT: FORMAT

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ANSWER 7 OF 7 HCAPLUS COPYRIGHT 2011 ACS on STN (Continued) Title compds. (I; X=SO2, POR10; Y=5-10 membered heteroaryl, Ph, naphthyl; Z=O, NH, CH2, S; RS=H, alkyl; RS, R7=H, Me; R8=H,
H, alkyl, cycloalkyl, heteroaryl, heterocycloalkyl; R11R12 = atoms to
form
5-10 membered mono- or bicyclic (heterocyclic) ring, Ph, naphthyl; dotted
line = optional double bond), were prepared Thus,
(1R,2S)-2=[[[4-(2-butynyloxy)phenyl]sulfonyl]methylamino]-N-
hydroxycyclopentanecarboxamide [preparation from
cis-2-amino-1-cyclopentanecarboxylic acid and
4-(2-butynyloxy)phenylsulfonyl chloride given] inhibited TACE with IC50 =
14 nM.

OS.CITING REF COUNT: 4 THERE ARE 4 CAPLUS RECORDS THAT CITE THIS
RECORD
                                                                     (4 CITINGS) THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE
REFERENCE COUNT:
FORMAT
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=> log y COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 52.24 519.45

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE TOTAL ENTRY SESSION

CA SUBSCRIBER PRICE

-8.70 -15.66

STN INTERNATIONAL LOGOFF AT 09:58:14 ON 22 FEB 2011